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CLAIMS

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- 1. A process for preparing carbohydrate fatty-acid esters comprising the steps of:
 - (a) reacting acylated carbohydrate with free fatty acid in the presence of an acid catalyst, under reduced pressure;
 - (b) decolorizing and separating out the unreacted fatty acid, from the reaction mixture obtained in step (a);
- (c) precipitating out the unreacted acylated carbohydrate from the reaction mixture obtained in step (b); and
 - (d) recovering carbohydrate fatty ester from the reaction mixture obtained in step (c).
- 2. The process of preparing carbohydrate fatty-acid esters of Claim 1, wherein in step (a), no solvent is added thereto.
 - 3. The process of preparing carbohydrate fatty-acid esters of claim 1, wherein in the unreacted fatty-acid in the reaction mixture in step (b) is removed by precipitation from a solvent mixture at controlled temperature.
 - 4. The process of preparing carbohydrate fatty-acid esters of claim 1, wherein the unreacted fatty-acid in the reaction mixture in step (b) is removed from the reaction mixture by solvent extraction.

5. The process of preparing carbohydrate fatty acid ester of claim 1 wherein the unreacted acylated carbohydrate is precipitated out in step (c) by cooling the reaction mixture in step (b) to a temperature in the range of –4 to 10 degree C.

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6. The process of preparing carbohydrate fatty acid esters of claim 1, wherein the unreacted free fatty acids and the unreacted C2 or C3-acylated carbohydrate esters which are removed during purification steps (b) and (c) are recycled to the reactant mixture.

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7. The process of preparing carbohydrate fatty-acid ester of claim 1 wherein step (a) is carried out at a pressure in the range of 4 - 20 Torr.

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8. The process of preparing carbohydrate fatty-acid ester of claim 1 wherein step (a) is carried out at a pressure in the range of 5-10 Torr.

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9. The process of preparing carbohydrate fatty-acid esters of Claim 1, wherein mono-, di- and poly-fatty acid esters of C2- or C3-acylated carbohydrates of various HLB values are obtained.

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10. The process of preparing carbohydrate fatty-acid esters of Claim 1, wherein the HLB values of the product carbohydrate fatty-acid esters are in the range of 1 to 10.

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- 11 The process of preparing carbohydrate fatty-acid esters of Claim 1, further comprising the steps of:-
- (e) librating free hydroxyl groups by partial hydrolysis of the C2- or C3-acylated carbohydrate fatty acid ester in the presence of an acid catalyst for a predetermined period of time to obtain carbohydrate fatty acid ester having free hydroxyl groups of predetermined HLB values.
- 12. The process of preparing carbohydrate fatty acid esters of claim 11, wherein the HLB values of the product carbohydrate fatty-acid esters are in the range of 8 to 16.
- 13. The process of preparing carbohydrate fatty-acid esters of Claim 1, wherein step (a) is processed at a temperature ranging from 60 to 95 degree C.
- 14. A process of preparing carbohydrate fatty acid esters comprising the steps of:
- (a) reacting acylated carbohydrate with free fatty acid in the presence of an acid catalyst, under reduced pressure;
 - (b) decolorizing and separating out the unreacted fatty acid, from the reaction mixture obtained in step (a);
 - (c) precipitating out the unreacted acylated carbohydrate from the reaction mixture obtained in step (a);
- 25 (d) removing the unreacted free fatty acids and carbohydrate esters of low molecular-weight carboxylic acids during purification,

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and recycling the removed unreacted free fatty acids and carbohydrate esters to the starting reactant mixture; and

- (e) librating free hydroxyl groups by partial hydrolysis of the acylated carbohydrate fatty acid ester in the presence of an acid catalyst for a predetermined period of time to obtain carbohydrate fatty acid ester having free hydroxyl groups of predetermined HLB values.
- 15. Carbohydrate fatty-acid esters produced in accordance with the process of claim 1 or 14.
- 16. The process according to claims 1 or 14 wherein the reactant carbohydrates include the group consisting of partially or peracylated mono-, di- and tri-saccharides in which the monosaccharide units could be a furanosyl, pyranosyl or a C2-C6 open-chain structure.
- 17. The process according to claims 1, 14 or 16 wherein the acyl group in the reactant acylated carbohydrates is acetic or propanoic acyl group.
- 18. The process according to claims 1 or 14 wherein, the acid catalysts includes sulphuric and camphorsulfonic acids, in the case of the monosaccharides; and boron trifluoride diethyl etherate, alkyl sulphonic acid polysiloxanes and tosylic acid, in the case of the diand tri-saccharides.

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- 19. The process according to claims 1 or 14 wherein the workup solvents includes ethanol, iso-propanol, n-propanol and ethyl acetate.
- 20. The process according to claims 4 wherein the extraction solvent is hexane
 - 21. The process according to claims 1 or 14 wherein the reactant free fatty acids have C6–C22 chain-length, with zero, mono or diunsaturations.
 - 22. The process according to claims 11 or 14 wherein the hydrolysis acid catalyst is trifluoroacetic acid.
 - 23. The process according to claims 11 or 14 wherein the partially hydrolysed carbohydrate fatty acid esters are further separated by stage cooling, at controlled temperature ranging from -15 to 10 degree C, according to their degree of acylation.